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SENATE BILL 272

43RD LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 1997

INTRODUCED BY

MICHAEL S. SANCHEZ

AN ACT

RELATING TO LICENSURE; CLARIFYING THE PRACTICE OF ORIENTAL
MEDICINE; GIVING DOCTORS OF ORIENTAL MEDICINE PRESCRIPTIVE
AUTHORITY; DESIGNATING DOCTORS OF ORIENTAL MEDICINE AS PRIMARY
CARE PROVIDERS; PROVIDING FOR ANNUAL LICENSURE; INCREASING FEES;
EXPANDING THE AUTHORITY TO DENY, SUSPEND OR REVOKE A LICENSE;
REQUIRING LICENSEES TO PAY COSTS OF DISCIPLINARY PROCEEDINGS
UNDER CERTAIN CIRCUMSTANCES.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

Section 1. Section 26-1-2 NMSA 1978 (being Laws 1967,
Chapter 23, Section 2, as amended) is amended to read:

"26-1-2. DEFINITIONS. --As used in the New Mexico Drug,
Device and Cosmetic Act:

A. "board" means the board of pharmacy or its duly
authorized agent;

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1 B. "person" includes individual, partnership,
2 corporation, association, institution or establishment;

3 C. "biological product" means any virus, therapeutic
4 serum, toxin, antitoxin or analogous product applicable to the
5 prevention, treatment or cure of diseases or injuries of man and
6 domestic animals and, as used within the meaning of this
7 definition:

8 (1) a "virus" is interpreted to be a product
9 containing the minute living cause of an infectious disease and
10 includes [~~but is not limited to~~] filterable viruses, bacteria,
11 rickettsia, fungi and protozoa;

12 (2) a "therapeutic serum" is a product obtained
13 from blood by removing the clot or clot components and the blood
14 cells;

15 (3) a "toxin" is a product containing a soluble
16 substance poisonous to laboratory animals or man in doses of one
17 milliliter or less of the product and having the property,
18 following the injection of nonfatal doses into an animal, or
19 causing to be produced therein another soluble substance [~~which~~]
20 that specifically neutralizes the poisonous substance and
21 [~~which~~] that is demonstrable in the serum of the animal thus
22 immunized; and

23 (4) an "antitoxin" is a product containing the
24 soluble substance in serum or other body fluid of an immunized
25 animal [~~which~~] that specifically neutralizes the toxin against

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1 which the animal is immune;

2 D. "controlled substance" means any drug, substance
3 or immediate precursor enumerated in Schedules I through V of
4 the Controlled Substances Act;

5 E. "drug" means:

6 (1) articles recognized in an official
7 compendium;

8 (2) articles intended for use in the diagnosis,
9 cure, mitigation, treatment or prevention of disease in man or
10 other animals and includes the domestic animal biological
11 products regulated under the federal Virus-Serum-Toxin Act, 37
12 Stat 832-833, 21 U.S.C. 151-158 and the biological products
13 applicable to man regulated under Federal 58 Stat 690, as
14 amended, 42 U.S.C. 216, Section 351, and 58 Stat 702, as
15 amended, 42 U.S.C. 262;

16 (3) articles other than food [~~which~~] that
17 affect the structure or any function of the body of man or other
18 animals; and

19 (4) articles intended for use as a component of
20 Paragraph (1), (2) or (3) of this subsection, but does not
21 include devices or their component parts or accessories;

22 F. "dangerous drug" means a drug, other than a
23 controlled substance enumerated in Schedule I of the Controlled
24 Substances Act, [~~which~~] that because of any potentiality for
25 harmful effect or the method of its use or the collateral

1 measures necessary to its use is not safe except under the
2 supervision of a practitioner licensed by law to direct the use
3 of such drug and hence for which adequate directions for use
4 cannot be prepared. "Adequate directions for use" means
5 directions under which the layman can use a drug or device
6 safely and for the purposes for which it is intended. A drug
7 shall be dispensed only upon the prescription of a practitioner
8 licensed by law to administer or prescribe such drug if it:

9 (1) is a habit-forming drug and contains any
10 quantity of a narcotic or hypnotic substance, or any chemical
11 derivative of such substance, [~~which~~] that has been found under
12 the federal act and the board to be habit-forming;

13 (2) because of its toxicity or other
14 potentiality for harmful effect or the method of its use or the
15 collateral measures necessary to its use is not safe for use
16 except under the supervision of a practitioner licensed by law
17 to administer or prescribe such drug;

18 (3) is limited by an approved application by
19 Section 505 of the federal act to the use under the professional
20 supervision of a practitioner licensed by law to administer or
21 prescribe such drug;

22 (4) bears the legend: "Caution: federal law
23 prohibits dispensing without prescription."; or

24 (5) bears the legend: "Caution: federal law
25 restricts this drug to use by or on the order of a licensed

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1 veterinarian. ";

2 G. "counterfeit drug" means a drug other than a
3 controlled substance [~~which~~] that, or the container or labeling
4 of which, without authorization, bears the trademark, trade name
5 or other identifying mark, imprint or device, or any likeness,
6 of a drug manufacturer, processor, packer or distributor other
7 than the person who in fact manufactured, processed, packed or
8 distributed such drug and [~~which~~] that falsely purports or is
9 represented to be the product of or to have been packed or
10 distributed by such other drug manufacturer, processor, packer
11 or distributor;

12 H. "device", except when used in Subsection P of
13 this section and in Subsection G of Section 26-1-3, Subsection L
14 and Paragraph (4) of Subsection A of Section 26-1-11 and
15 Subsection C of Section 26-1-24 NMSA 1978, means an instrument,
16 apparatus, implement, machine, contrivance, implant, in vitro
17 reagent or other similar or related article, including any
18 component, part or accessory, [~~which~~] that is:

- 19 (1) recognized in an official compendium;
- 20 (2) intended for use in the diagnosis of
21 disease or other conditions, or in the cure, mitigation,
22 treatment or prevention of disease, in man or other animals; or
- 23 (3) intended to affect the structure or any
24 function of the body of man or other animals and [~~which~~] that
25 does not achieve any of its principal intended purposes through

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1 chemical action within or on the body of man or other animals
2 and which is not dependent upon being metabolized for
3 achievement of any of its principal intended purposes;

4 I. "prescription" means an order given individually
5 for the person for whom prescribed, either directly from the
6 prescriber to the pharmacist or indirectly by means of a written
7 order signed by the prescriber, and bearing the name and address
8 of the prescriber, his license classification, the name and
9 address of the patient, the name and quantity of the drug
10 prescribed, directions for use and the date of issue. No person
11 other than a practitioner shall prescribe or write a
12 prescription;

13 J. "practitioner" means a physician, doctor of
14 oriental medicine, dentist, veterinarian or other person
15 licensed to prescribe and administer drugs [~~which~~] that are
16 subject to the New Mexico Drug, Device and Cosmetic Act;

17 K. "cosmetic" means:

18 (1) articles intended to be rubbed, poured,
19 sprinkled or sprayed on, introduced into or otherwise applied to
20 the human body or any part thereof for cleansing, beautifying,
21 promoting attractiveness or altering the appearance; and

22 (2) articles intended for use as a component of
23 any articles enumerated in Paragraph (1) of this subsection,
24 except that the term shall not include soap;

25 L. "official compendium" means the official United

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1 States pharmacopoeia national formulary or the official
2 homeopathic pharmacopoeia of the United States or any supplement
3 to either of them;

4 M. "label" means a display of written, printed or
5 graphic matter upon the immediate container of any article. A
6 requirement made by or under the authority of the New Mexico
7 Drug, Device and Cosmetic Act that any word, statement or other
8 information appear on the label shall not be considered to be
9 complied with unless the word, statement or other information
10 also appears on the outside container or wrapper, if any, of the
11 retail package of the article or is easily legible through the
12 outside container or wrapper;

13 N. "immediate container" does not include package
14 liners;

15 O. "labeling" means all labels and other written,
16 printed or graphic matter:

17 (1) upon any article or any of its containers
18 or wrappers; or

19 (2) accompanying any article;

20 P. "misbranded" means a label to an article [~~which~~]
21 that is misleading. In determining whether the label is
22 misleading, there shall be taken into account, among other
23 things, not only representations made or suggested by statement,
24 word, design, device or any combination of the foregoing, but
25 also the extent to which the label fails to reveal facts

1 material in the light of such representations or material with
2 respect to consequences [~~which~~] that may result from the use of
3 the article to which the label relates under the conditions of
4 use prescribed in the label or under such conditions of use as
5 are customary or usual;

6 Q. "advertisement" means all representations
7 disseminated in any manner or by any means, other than by
8 labeling, for the purpose of inducing, or [~~which~~] that are
9 likely to induce, directly or indirectly, the purchase of drugs,
10 devices or cosmetics;

11 R. "antiseptic", when used in the labeling or
12 advertisement of an antiseptic, shall be considered to be a
13 representation that it is a germicide, except in the case of a
14 drug purporting to be or represented as an antiseptic for
15 inhibitory use as a wet dressing, ointment, dusting powder or
16 such other use as involves prolonged contact with the body;

17 S. "new drug" means:

18 (1) any drug, the composition of which is such
19 that the drug is not generally recognized, among experts
20 qualified by scientific training and experience to evaluate the
21 safety and efficacy of drugs, as safe and effective for use
22 under the conditions prescribed, recommended or suggested in the
23 labeling thereof; or

24 (2) any drug, the composition of which is such
25 that the drug, as a result of investigation to determine its

1 safety and efficacy for use under such conditions, has become so
2 recognized, but [~~which~~] that has not, otherwise than in such
3 investigations, been used to a material extent or for a material
4 time under such conditions;

5 T. "contaminated with filth" applies to any drug,
6 device or cosmetic not securely protected from dirt, dust and,
7 as far as may be necessary by all reasonable means, from all
8 foreign or injurious contaminations, or any drug, device or
9 cosmetic found to contain any dirt, dust, foreign or injurious
10 contamination or infestation;

11 U. "selling of drugs, devices or cosmetics" shall be
12 considered to include the manufacture, production, processing,
13 packing, exposure, offer, possession and holding of any such
14 article for sale and the sale and the supplying or applying of
15 any such article in the conduct of any drug or cosmetic
16 establishment;

17 V. "color additive" means a material [~~which~~] that:

18 (1) is a dye, pigment or other substance made
19 by a process of synthesis or similar artifice or extracted,
20 isolated or otherwise derived, with or without intermediate or
21 final change of identity, from a vegetable, mineral, animal or
22 other source; or

23 (2) when added or applied to a drug or cosmetic
24 or to the human body or any part thereof, is capable, alone or
25 through reaction with other substances, of imparting color

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1 thereto; except that such term does not include any material
2 [~~which~~] that has been or hereafter is exempted under the federal
3 act;

4 W. "federal act" means the Federal Food, Drug and
5 Cosmetic Act;

6 X. "restricted device" means a device for which the
7 sale, distribution or use is lawful only upon the written or
8 oral authorization of a practitioner licensed by law to
9 administer, prescribe or use the device and for which the
10 federal food and drug administration requires special training
11 or skills of the practitioner to use or prescribe. This
12 definition does not include custom devices defined in the
13 federal act and exempt from performance standards or premarket
14 approval requirements under Section 520 (b) of the federal act;
15 and

16 Y. "prescription device" means a device [~~which~~]
17 that, because of its potential for harm, the method of its use
18 or the collateral measures necessary to its use, is not safe
19 except under the supervision of a practitioner licensed in this
20 state to direct the use of such device and for which "adequate
21 directions for use" cannot be prepared, but that bears the
22 label: "Caution: Federal law restricts this device to sale by
23 or on the order of a _____", the blank to be filled with
24 the word "physician", "doctor of oriental medicine", "dentist",
25 "veterinarian" or with the descriptive designation of any other

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1 practitioner licensed in this state to use or order the use of
2 the device. "

3 Section 2. Section 61-14A-1 NMSA 1978 (being Laws 1993,
4 Chapter 158, Section 9) is amended to read:

5 "61-14A-1. SHORT TITLE. -- [~~Sections 61-14A-1 through~~
6 ~~61-14A-24~~] Chapter 61, Article 14A NMSA 1978 may be cited as the
7 "Acupuncture and Oriental Medicine Practice Act". "

8 Section 3. Section 61-14A-3 NMSA 1978 (being Laws 1993,
9 Chapter 158, Section 11) is amended to read:

10 "61-14A-3. DEFINITIONS. -- As used in the Acupuncture and
11 Oriental Medicine Practice Act:

12 A. "acupuncture" means the use of needles inserted
13 into and removed from the human body and the use of other
14 devices, modalities and procedures at specific locations on the
15 body for the prevention, cure or correction of any disease,
16 illness, injury, pain or other condition by controlling and
17 regulating the flow and balance of energy and functioning of the
18 person to restore and maintain health;

19 B. "board" means the board of acupuncture and
20 oriental medicine;

21 [~~C. "department" means the regulation and licensing~~
22 ~~department;~~

23 ~~D.]~~ C. "doctor of oriental medicine" means a
24 [~~physician~~] person licensed as a physician to practice
25 acupuncture and oriental medicine [~~and includes the terms~~

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1 ~~"oriental medical physician", "doctor of acupuncture",~~
2 ~~"acupuncture physician", "acupuncture practitioner" and~~
3 ~~"acupuncturist"]~~ with the ability to practice independently.
4 serve as a primary care provider and as necessary collaborate
5 with other health care providers;

6 [E.] D. "moxibustion" means the use of heat on or
7 above specific locations or on acupuncture needles at specific
8 locations on the body for the prevention, cure or correction of
9 any disease, illness, injury, pain or other condition;

10 [F.] E. "oriental medicine" means the distinct
11 system of primary health care that uses all allied techniques of
12 oriental medicine, both traditional and modern, to diagnose,
13 treat and prescribe ~~[as defined in Subsection 6 of this section]~~
14 for the prevention, cure or correction of any disease, illness,
15 injury, pain or other physical or mental condition by
16 controlling and regulating the flow and balance of energy and
17 functioning of the person to restore and maintain health; ~~and]~~

18 F. "primary care provider" means a health care
19 professional who provides the first level of basic or general
20 health care for an individual's health needs, including
21 diagnostic and treatment services; and

22 G. "techniques of oriental medicine" means:

23 (1) the diagnostic and treatment techniques
24 ~~[utilized]~~ used in oriental medicine that include ~~[but are not~~
25 ~~limited to]~~ diagnostic procedures; acupuncture; moxibustion;

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1 manual therapy, also known as tui na; other physical medicine
2 modalities and therapeutic procedures; breathing and exercise
3 techniques; and dietary, nutritional and lifestyle counseling;
4 [~~and~~]

5 (2) the prescription or administration of any
6 herbal medicine, homeopathic medicine [~~vitamin, mineral, enzyme,~~
7 ~~glandular or nutritional supplement~~] or other substances,
8 including vitamins, minerals, enzymes, glandular products, amino
9 acids, dietary and nutritional supplements; and

10 (3) the prescription or administration of
11 biological products, drugs, dangerous drugs and cosmetics, other
12 than those enumerated in Paragraph (2) of this subsection, and
13 the prescription or administration of devices, restricted
14 devices and prescription devices, as these substances and
15 devices are defined in the New Mexico Drug, Device and Cosmetic
16 Act, if the board determines by rule that any such substance or
17 device is necessary in the practice of oriental medicine. "

18 Section 4. Section 61-14A-5 NMSA 1978 (being Laws 1993,
19 Chapter 158, Section 13) is amended to read:

20 "61-14A-5. TITLE. -- Any person licensed [~~under~~] pursuant to
21 provisions of the Acupuncture and Oriental Medicine Practice
22 Act, in advertising his services to the public, shall use the
23 title "doctor of oriental medicine" or "D. O. M ". [~~Effective~~
24 ~~July 1, 1994~~] The title "doctor of oriental medicine" or
25 "D. O. M " shall supersede the use of all other titles that

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1 include the words "medical doctor" or the initials "M D." unless
2 the person is a medical doctor licensed pursuant to provisions
3 of the Medical Practice Act. "

4 Section 5. Section 61-14A-6 NMSA 1978 (being Laws 1993,
5 Chapter 158, Section 14) is amended to read:

6 "61-14A-6. EXEMPTIONS. --

7 ~~[A. Nothing in the Acupuncture and Oriental Medicine~~
8 ~~Practice Act is intended to limit, interfere with or prevent any~~
9 ~~other class of licensed health care professionals from~~
10 ~~practicing within the scope of their license as defined by each~~
11 ~~profession's New Mexico licensing statutes, but they shall not~~
12 ~~hold themselves out to the public or any private group or~~
13 ~~business by using any title or description of services that~~
14 ~~includes the terms acupuncture, acupuncturist or oriental~~
15 ~~medicine unless they are licensed under the Acupuncture and~~
16 ~~Oriental Medicine Practice Act.~~

17 B.] A. Students enrolled in an educational program
18 in acupuncture and oriental medicine approved by the board may
19 practice acupuncture and oriental medicine under the direct
20 supervision of a teacher at an institute or with a private tutor
21 as part of the educational program in which they are enrolled.

22 ~~[C.]~~ B. The Acupuncture and Oriental Medicine
23 Practice Act shall not apply to or affect the following
24 practices [~~provided that~~] if the individual does not hold
25 himself out as a doctor of oriental medicine or as practicing

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1 acupuncture or oriental medicine:

2 (1) the administering of gratuitous services in
3 cases of emergency;

4 (2) the domestic administering of family
5 remedies;

6 (3) the counseling about or the teaching and
7 demonstration of breathing and exercise techniques;

8 (4) the counseling or teaching about diet and
9 nutrition;

10 (5) the spiritual or lifestyle counseling of
11 any individual or spiritual group or the practice of the
12 religious tenets of any church; [øæ]

13 (6) the providing of information about the
14 general usage of herbal medicines, homeopathic medicines,
15 vitamins, minerals, enzymes or glandular or nutritional
16 supplements; or

17 (7) the use of needles for diagnostic purposes
18 and the use of needles for the administration of diagnostic or
19 therapeutic substances by licensed health care professionals."

20 Section 6. Section 61-14A-10 NMSA 1978 (being Laws 1993,
21 Chapter 158, Section 18) is amended to read:

22 "61-14A-10. REQUIREMENTS FOR LICENSING. --The board shall
23 grant a license to practice acupuncture and oriental medicine to
24 any person who has submitted to the board:

25 A. the completed application for licensing on the

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1 form provided by the board;

2 B. the required documentation as determined by the
3 board;

4 C. the required fees;

5 D. an affidavit stating that the applicant has not
6 been found guilty of unprofessional conduct or incompetency;

7 E. proof, as determined by the board, that the
8 applicant has completed [~~an~~] a board-approved educational
9 program in acupuncture and oriental medicine as provided for in
10 the Acupuncture and Oriental Medicine Practice Act and the rules
11 [~~and regulations~~] of the board; and

12 F. proof that he has passed [~~an examination~~] the
13 examinations approved by the board. "

14 Section 7. Section 61-14A-11 NMSA 1978 (being Laws 1993,
15 Chapter 158, Section 19) is amended to read:

16 "61-14A-11. EXAMINATIONS. --

17 A. The board shall establish procedures to ensure
18 that examinations for licensing are offered at least once a
19 year.

20 B. The board shall establish by rule the deadline
21 for receipt of the application for licensing examination and
22 other rules relating to the taking and retaking of licensing
23 examinations.

24 C. The board shall establish by rule the passing
25 grades for its approved examinations.

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1 D. The board ~~may~~ approve by rule examinations that
2 are used for national certification or other examinations.

3 E. The board shall require each qualified applicant
4 to pass a written examination that includes, as a minimum, the
5 following subjects:

6 (1) anatomy and physiology;

7 (2) pathology;

8 (3) diagnosis; [~~and~~]

9 (4) pharmacology; and

10 [~~(4)~~] (5) principles, practices and treatment
11 techniques of acupuncture and oriental medicine.

12 F. The board [~~shall~~] may require each qualified
13 applicant to pass a practical examination that demonstrates his
14 knowledge of and skill in the application of the diagnostic and
15 treatment techniques of acupuncture and oriental medicine.

16 G. The board shall require each qualified applicant
17 to pass a written or a practical examination or both in the
18 following subjects:

19 (1) hygiene, sanitation and clean-needle
20 technique; and

21 (2) needle and instrument sterilization
22 techniques.

23 H. The board ~~may~~ require each qualified applicant to
24 pass a written examination on the state laws and [~~regulations~~]
25 rules that pertain to the practice of acupuncture and oriental

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1 medicine. "

2 Section 8. Section 61-14A-13 NMSA 1978 (being Laws 1993,
3 Chapter 158, Section 21, as amended) is amended to read:

4 "61-14A-13. REQUIREMENTS FOR RECIPROCAL LICENSING. --The
5 board may grant a license to practice acupuncture and oriental
6 medicine to a person who has been licensed, certified,
7 registered or legally recognized as a doctor of oriental
8 medicine in another state, district or territory of the United
9 States or foreign country if the applicant:

10 A. submits the completed application for reciprocal
11 licensing on the form provided by the board;

12 B. submits the required documentation as determined
13 by the board;

14 C. submits the required fee for application for
15 reciprocal licensing;

16 D. submits an affidavit stating that the applicant
17 has not been found guilty of unprofessional conduct or
18 incompetency;

19 E. has passed a practical examination that
20 demonstrates his knowledge of and skill in the application of
21 the diagnostic and treatment techniques of acupuncture and
22 oriental medicine, if the board requires regular applicants to
23 pass a practical examination, or within the last six years has
24 five years of clinical experience, as defined by rule, in the
25 practice of acupuncture and oriental medicine;

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1 F. has passed a written examination on the state
2 laws and rules that pertain to the practice of acupuncture and
3 oriental medicine, if the board requires regular applicants for
4 licensure to pass such an examination;

5 [F.] G. is licensed, certified, registered or
6 legally recognized as a doctor of oriental medicine in another
7 state, district or territory of the United States or foreign
8 country in which the requirements for practice are similar to
9 those of this state; and

10 [G.] H. is licensed, certified, registered or
11 legally recognized as a doctor of oriental medicine in a state,
12 district or territory of the United States or foreign country
13 that permits a doctor of oriental medicine licensed under the
14 provisions of the Acupuncture and Oriental Medicine Practice Act
15 to practice acupuncture and oriental medicine in that
16 jurisdiction by reciprocal credentials review. "

17 Section 9. Section 61-14A-14 NMSA 1978 (being Laws 1993,
18 Chapter 158, Section 22) is amended to read:

19 "61-14A-14. APPROVAL OF EDUCATIONAL PROGRAMS. --

20 A. The board shall establish by rule the criteria
21 for board approval of educational programs in acupuncture and
22 oriental medicine. For [the] an educational program in
23 acupuncture and oriental medicine to meet board approval, proof
24 shall be submitted to the board demonstrating that the
25 educational program:

1 (1) was for a period of not less than four
2 academic years;

3 (2) included a minimum of seven hundred fifty
4 hours of supervised clinical practice;

5 (3) was taught by qualified teachers or a
6 qualified private tutor;

7 (4) required as a prerequisite to graduation
8 personal attendance in all classes and clinics and, as a
9 minimum, the completion of the following subjects:

10 (a) anatomy and physiology;

11 (b) pathology;

12 (c) diagnosis;

13 (d) pharmacology;

14 [~~(d)~~] (e) oriental principles of life
15 therapy, including diet, nutrition and counseling;

16 [~~(e)~~] (f) theory and techniques of
17 traditional and modern acupuncture and oriental medicine;

18 [~~(f)~~] (g) precautions and
19 contraindications for acupuncture treatment;

20 [~~(g)~~] (h) theory and application of
21 meridian pulse evaluation and meridian point location;

22 [~~(h)~~] (i) traditional and modern methods
23 of life-energy evaluation;

24 [~~(i)~~] (j) the prescription of herbal
25 medicine and precautions and contraindications for its use;

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1 approval of an educational program.

2 C. Institutes and private tutors outside New Mexico
3 that offer educational programs in acupuncture and oriental
4 medicine with the intent to graduate students qualified to be
5 applicants for licensing examination by the board may have their
6 educational programs annually approved by the board. For the
7 educational program in acupuncture and oriental medicine to be
8 approved by the board, the institute or private tutor shall
9 submit:

10 (1) the completed application for approval of
11 an educational program;

12 (2) the required documentation as determined by
13 the board;

14 (3) proof, as determined by the board, that the
15 educational requirements [~~referred to~~] provided for in
16 Subsection A of this section are being met; and

17 (4) the required fee for application for
18 approval of an educational program.

19 D. Each institute and private tutor in New Mexico
20 that offers an approved educational program in acupuncture and
21 oriental medicine as referred to in Subsection B of this section
22 shall renew their approval annually by submitting:

23 (1) the completed application for renewal of
24 approval of an educational program on the form provided by the
25 board;

1 (2) proof, as determined by the board, that the
2 educational requirements [~~referred to~~] provided for in
3 Subsection A of this section are being met; and

4 (3) the required fee for application for
5 renewal of approval of an educational program.

6 E. Each institute and private tutor outside New
7 Mexico that offers an approved educational program in
8 acupuncture and oriental medicine as referred to in Subsection C
9 of this section may renew their approval annually by submitting:

10 (1) the completed application for renewal of
11 approval of an educational program on the form provided by the
12 board;

13 (2) proof, as determined by the board, that the
14 educational requirements [~~referred to~~] provided for in
15 Subsection A of this section are being met; and

16 (3) the required fee for application for
17 renewal of approval of an educational program.

18 F. A sixty-day grace period shall be allowed each
19 institute or private tutor after the end of the approval period,
20 during which time the approval may be renewed by submitting:

21 (1) the completed application for renewal of
22 approval of an educational program on the form provided by the
23 board;

24 (2) proof, as determined by the board, that the
25 educational requirements [~~referred to~~] provided for in

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1 Subsection A of this section are being met;

2 (3) the required fee for application for
3 renewal of approval of an educational program; and

4 (4) the required fee for late renewal of
5 approval.

6 G. Any approval not renewed at the end of the grace
7 period shall be considered expired. For renewal of an expired
8 approval, the board shall establish by rule any requirements or
9 fees that are in addition to the fee for annual renewal of
10 approval and may require the institute or private tutor to
11 reapply as a new applicant. "

12 Section 10. Section 61-14A-15 NMSA 1978 (being Laws 1993,
13 Chapter 158, Section 23) is amended to read:

14 "61-14A-15. LICENSE RENEWAL. --

15 A. Each licensee shall renew his license
16 [~~biennially~~] annually by submitting:

17 (1) the ~~completed~~ application for license
18 renewal on the form provided by the board; and

19 (2) the required fee for [~~biennial~~] annual
20 license renewal.

21 B. The board may require proof of continuing
22 education or other proof of competency as a requirement for
23 renewal.

24 C. A sixty-day grace period shall be allowed each
25 licensee after the end of the licensing period, during which

Underscored material = new
[bracketed material] = delete

1 time the license may be renewed by submitting:

2 (1) the completed application for license
3 renewal on the form provided by the board;

4 (2) the required fee for [~~biennial~~] annual
5 license renewal; and

6 (3) the required fee for late license renewal.

7 D. Any license not renewed at the end of the grace
8 period shall be considered expired and the licensee shall not be
9 eligible to practice within the state. For renewal of an
10 expired license, the board shall establish by rule any
11 requirements or fees that are in addition to the fee for
12 [~~biennial~~] annual license renewal and may require the former
13 licensee to reapply as a new applicant. "

14 Section 11. Section 61-14A-16 NMSA 1978 (being Laws 1993,
15 Chapter 158, Section 24) is amended to read:

16 "61-14A-16. FEES. --The board shall establish a schedule of
17 reasonable nonrefundable fees not to exceed the following
18 amounts:

- 19 A. application for licensing \$[~~500~~] 1,000;
- 20 B. application for reciprocal licensing [~~750~~] 1,500;
- 21 C. application for temporary licensing . [~~300~~] 600;
- 22 D. examination, not including the cost of any
23 nationally recognized examination [~~350~~] 1,500;
- 24 E. [~~biennial~~] annual license renewal 400;
- 25 F. late license renewal [~~200~~] 400;

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- 1 G. expired license renewal [~~400~~] 800;
- 2 H. temporary license renewal [~~100~~] 200;
- 3 I. application for approval or renewal of approval
- 4 of an educational program [~~400~~] 800;
- 5 J. late renewal of approval of an educational
- 6 program [~~200~~] 400;
- 7 K. expired renewal of approval of an educational
- 8 program [~~400~~] 800;
- 9 L. annual continuing education provider
- 10 registration [~~200~~;
- 11 and] 400;

12 M duplicate license \$100; and

13 [~~M-~~] N. any and all fees to cover reasonable and
14 necessary administrative expenses. "

15 Section 12. Section 61-14A-17 NMSA 1978 (being Laws 1993,
16 Chapter 158, Section 25) is amended to read:

17 "61-14A-17. DISCIPLINARY PROCEEDINGS-- JUDICIAL REVIEW -
18 APPLICATION OF UNIFORM LICENSING ACT. --

19 A. In accordance with the procedures contained in
20 the Uniform Licensing Act, the board may deny, revoke or suspend
21 any permanent or temporary license held or applied for under the
22 Acupuncture and Oriental Medicine Practice Act, upon findings by
23 the board that the licensee or applicant:

24 (1) is guilty of fraud or deceit in procuring
25 or attempting to procure a license;

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1 (2) has been convicted of a felony. A
2 certified copy of the record of conviction shall be conclusive
3 evidence of such conviction;

4 (3) is guilty of incompetence as defined by
5 board rule;

6 (4) is habitually intemperate, is addicted to
7 the use of habit-forming drugs or is addicted to any vice to
8 such a degree as to render him unfit to practice as a doctor of
9 oriental medicine;

10 (5) is guilty of unprofessional conduct, as
11 defined by board rule;

12 (6) is guilty of any violation of the
13 Controlled Substances Act;

14 (7) has violated any provision of the
15 Acupuncture and Oriental Medicine Practice Act or rules [~~and~~
16 ~~regulations adopted~~] promulgated by the board;

17 (8) is guilty of failing to furnish the board,
18 its investigators or representatives with information requested
19 by the board;

20 (9) is guilty of willfully or negligently
21 practicing beyond the scope of acupuncture and oriental medicine
22 as defined in the Acupuncture and Oriental Medicine Practice
23 Act;

24 (10) is guilty of failing to adequately
25 supervise a sponsored temporary licensee;

Underscored material = new
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1 (11) is guilty of aiding or abetting the
2 practice of acupuncture and oriental medicine by a person not
3 licensed by the board;

4 (12) is guilty of practicing or attempting to
5 practice under an assumed name;

6 (13) advertises by means of knowingly false
7 statements;

8 (14) advertises or attempts to attract
9 patronage in any unethical manner prohibited by the Acupuncture
10 and Oriental Medicine Practice Act or the rules [~~and~~
11 ~~regulations~~] of the board;

12 (15) has been declared mentally incompetent by
13 regularly constituted authorities; [~~or~~]

14 (16) has had a license, certificate or
15 registration to practice as a doctor of oriental medicine
16 revoked, suspended or denied in any jurisdiction of the United
17 States or a foreign country for actions of the licensee similar
18 to acts described in this subsection. A certified copy of the
19 record of the jurisdiction taking such disciplinary action will
20 be conclusive evidence thereof; or

21 (17) fails, when diagnosing or treating a
22 patient, to possess or apply the knowledge or to use the skill
23 and care ordinarily used by reasonably well-qualified doctors of
24 oriental medicine practicing under similar circumstances, giving
25 due consideration to the locality involved.

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B. Disciplinary proceedings may be instituted by any person, shall be by sworn complaint and shall conform with the provisions of the Uniform Licensing Act. Any party to the hearing may obtain a copy of the hearing record upon payment of the costs of the copy.

C. Any person filing a sworn complaint shall be immune from liability arising out of civil action if the complaint is filed in good faith and without actual malice.

D. The licensee shall bear the costs of disciplinary proceedings unless exonerated."

FORTY-THIRD LEGISLATURE

FIRST SESSION, 1997

SB 272/a

February 14, 1997

Mr. President:

Your PUBLIC AFFAIRS COMMITTEE, to whom has been referred

SENATE BILL 272

has had it under consideration and reports same with recommendation that it DO PASS, amended as follows:

1. On page 12, line 19, after "professional" insert "acting within the scope of his license".

2. On page 13, line 17, before the period insert: "; provided, however, that for the purposes of the Acupuncture and Oriental Medicine Practice Act, "dangerous drug" does not include any controlled substance as defined in the Controlled Substances Act".

3. On page 14, lines 7 through 16, remove the beginning bracket and line-through.

FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

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SPAC/SB 272

Page 31

4. On page 14, line 17, remove the bracket and line-through and strike the underscored "A."

5. On page 14, line 22, remove the brackets and line-through and strike the underscored "B.", and thence referred to the JUDICIARY COMMITTEE.

Respectfully submitted,

Shannon Robinson, Chairman

Adopted _____ Not Adopted _____
(Chief Clerk) (Chief Clerk)

Date _____

The roll call vote was 4 For 1 Against

Yes: 4

No: Adair

Excused: Boitano, Ingle, Vernon, Smith

Absent: None

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FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

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SPAC/SB 272

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Page 32

. 116237. 1

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. 113836. 1

State of New Mexico
House of Representatives

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4 FORTY-THIRD LEGISLATURE
5 FIRST SESSION, 1997
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9 March 17, 1997
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11 Mr. Speaker:
12

13 Your CONSUMER AND PUBLIC AFFAIRS COMMITTEE, to
14 whom has been referred

15
16 SENATE BILL 272, as amended

17 has had it under consideration and reports same with
18 recommendation that it DO PASS, amended as follows:
19

20 1. On page 11, lines 21 and 22, remove the beginning bracket
21 and line-through.

22
23 2. On page 11, line 23, remove the line-through and end
24 bracket and strike the underscored "C. ".
25

3. Strike Senate Public Affairs Committee Amendment 2.

. 113836. 1

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FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

HCPAC/SB 272, a

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4. On page 13, line 7, after the bracket strike the remainder of the line, strike line 8 through "including" and insert in lieu thereof a comma.

5. On page 13, line 9, after the semicolon strike "and".

6. On page 13, strike lines 11 through 17 and insert in lieu thereof:

"devices, restricted devices and prescription devices, as those devices are defined in the New Mexico Drug, Device and Cosmetic Act, if the board determines by rule that such devices are necessary in the practice of oriental medicine and if the prescribing doctor of oriental medicine has fulfilled requirements for prescriptive authority in accordance with rules promulgated by the board for the devices enumerated in this paragraph;

(4) the prescription or administration of cosmetics, therapeutic serum and over-the-counter drugs, other than those enumerated in Paragraph (2) of this subsection, as those are defined in the New Mexico Drug, Device and Cosmetic Act, if the prescribing doctor of oriental medicine has fulfilled the requirements for prescriptive authority in accordance with rules promulgated by the board for the substances enumerated in this paragraph; and

(5) the prescription or administration of the following dangerous drugs as they are defined in the New Mexico

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FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

HCPAC/SB 272, a

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Drug, Device and Cosmetic Act, if the prescribing doctor of oriental medicine has fulfilled the requirements for prescriptive authority in accordance with rules promulgated by the board for the substances enumerated in this paragraph:

(a) vapocoolants;

(b) topical application of naturally occurring hormones; and

(c) any of the drugs or substances enumerated in Paragraphs (2) and (4) of this subsection if at any time these substances or drugs are classified as dangerous drugs. "".

7. On pages 24 through 26, strike Sections 10 and 11 in their entirety.

8. Renumber the succeeding section accordingly. ,

and thence referred to the APPROPRIATIONS AND FINANCE COMMITTEE.

FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

HCPAC/SB 272, a

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Respectfully submitted,

Gary King, Chairman

Adopted _____ Not Adopted _____
(Chief Clerk) (Chief Clerk)

Date _____

The roll call vote was 5 For 2 Against

Yes: 5

No: Crook, Dana

Excused: Johnson, Rios, Vigil

Absent: None

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Underscored material = new
[bracketed material] = delete

State of New Mexico House of Representatives

**FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997**

March 18, 1997

Mr. Speaker:

**Your APPROPRIATIONS AND FINANCE COMMITTEE, to
whom has been referred**

SENATE BILL 272, as amended

**has had it under consideration and reports same with
recommendation that it DO PASS.**

Respectfully submitted,

Max Coll, Chairman

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FORTY-THIRD LEGISLATURE
FIRST SESSION, 1997

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Adopted _____ Not Adopted _____
(Chief Clerk) (Chief Clerk)

Date _____

The roll call vote was 17 For 0 Against

Yes: 17

Excused: None

Absent: None

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Underscored material = new
~~[bracketed material] = delete~~